

09/ 930,782

Welcome to STN International! Enter x:x

LOGINID:ssspta1202txn

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

* * * * * Welcome to STN International * * * * *

NEWS 1 Web Page URLs for STN Seminar Schedule - N. America
NEWS 2 Apr 08 "Ask CAS" for self-help around the clock
NEWS 3 Apr 09 BEILSTEIN: Reload and Implementation of a New Subject Area
NEWS 4 Apr 09 ZDB will be removed from STN
NEWS 5 Apr 19 US Patent Applications available in IFICDB, IFIPAT, and IFIUDB
NEWS 6 Apr 22 Records from IP.com available in CAPLUS, HCAPLUS, and ZCAPLUS
NEWS 7 Apr 22 BIOSIS Gene Names now available in TOXCENTER
NEWS 8 Apr 22 Federal Research in Progress (FEDRIP) now available
NEWS 9 Jun 03 New e-mail delivery for search results now available
NEWS 10 Jun 10 MEDLINE Reload
NEWS 11 Jun 10 PCTFULL has been reloaded
NEWS 12 Jul 02 FOREGE no longer contains STANDARDS file segment
NEWS 13 Jul 22 USAN to be reloaded July 28, 2002;
saved answer sets no longer valid
NEWS 14 Jul 29 Enhanced polymer searching in REGISTRY
NEWS 15 Jul 30 NETFIRST to be removed from STN
NEWS 16 Aug 08 CANCERLIT reload
NEWS 17 Aug 08 PHARMAMarketLetter(PHARMAML) - new on STN
NEWS 18 Aug 08 NTIS has been reloaded and enhanced
NEWS 19 Aug 19 Aquatic Toxicity Information Retrieval (AQUIRE)
now available on STN
NEWS 20 Aug 19 IFIPAT, IFICDB, and IFIUDB have been reloaded
NEWS 21 Aug 19 The MEDLINE file segment of TOXCENTER has been reloaded
NEWS 22 Aug 26 Sequence searching in REGISTRY enhanced
NEWS 23 Sep 03 JAPIO has been reloaded and enhanced

NEWS EXPRESS February 1 CURRENT WINDOWS VERSION IS V6.0d,
CURRENT MACINTOSH VERSION IS V6.0a(ENG) AND V6.0Ja(JP),
AND CURRENT DISCOVER FILE IS DATED 05 FEBRUARY 2002
NEWS HOURS STN Operating Hours Plus Help Desk Availability
NEWS INTER General Internet Information
NEWS LOGIN Welcome Banner and News Items
NEWS PHONE Direct Dial and Telecommunication Network Access to STN
NEWS WWW CAS World Wide Web Site (general information)

Enter NEWS followed by the item number or name to see news on that specific topic.

All use of STN is subject to the provisions of the STN Customer agreement. Please note that this agreement limits use to scientific research. Use for software development or design or implementation of commercial gateways or other similar uses is prohibited and may result in loss of user privileges and other penalties.

* * * * * STN Columbus * * * * *

FILE 'HOME' ENTERED AT 11:23:42 ON 06 SEP 2002

09/ 930,782

=> file reg

COST IN U.S. DOLLARS

SINCE FILE

ENTRY

TOTAL

SESSION

FULL ESTIMATED COST

0.21

0.21

FILE 'REGISTRY' ENTERED AT 11:23:50 ON 06 SEP 2002

USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.

PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

COPYRIGHT (C) 2002 American Chemical Society (ACS)

STRUCTURE FILE UPDATES: 5 SEP 2002 HIGHEST RN 447396-35-2

DICTIONARY FILE UPDATES: 5 SEP 2002 HIGHEST RN 447396-35-2

TSCA INFORMATION NOW CURRENT THROUGH MAY 20, 2002

Please note that search-term pricing does apply when
conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Calculated physical property data is now available. See HELP PROPERTIES
for more information. See STNote 27, Searching Properties in the CAS
Registry File, for complete details:

<http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf>

=>

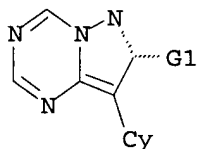
Uploading 09930782.str

L1 STRUCTURE UPLOADED

=> d l1

L1 HAS NO ANSWERS

L1 STR



G1 H,O

Structure attributes must be viewed using STN Express query preparation.

=> s l1

SAMPLE SEARCH INITIATED 11:24:07 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 494 TO ITERATE

100.0% PROCESSED 494 ITERATIONS

14 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 8547 TO 11213

PROJECTED ANSWERS: 56 TO 504

L2 14 SEA SSS SAM L1

=> s l1 ful

FULL SEARCH INITIATED 11:24:14 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 10201 TO ITERATE

09/ 930,782

100.0% PROCESSED 10201 ITERATIONS
SEARCH TIME: 00.00.02

324 ANSWERS

L3 324 SEA SSS FUL L1

=> file caplus

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

140.28

140.49

FILE 'CAPLUS' ENTERED AT 11:24:22 ON 06 SEP 2002

USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.

PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

COPYRIGHT (C) 2002 AMERICAN CHEMICAL SOCIETY (ACS)

Copyright of the articles to which records in this database refer is held by the publishers listed in the PUBLISHER (PB) field (available for records published or updated in Chemical Abstracts after December 26, 1996), unless otherwise indicated in the original publications. The CA Lexicon is the copyrighted intellectual property of the American Chemical Society and is provided to assist you in searching databases on STN. Any dissemination, distribution, copying, or storing of this information, without the prior written consent of CAS, is strictly prohibited.

FILE COVERS 1907 - 6 Sep 2002 VOL 137 ISS 11

FILE LAST UPDATED: 5 Sep 2002 (20020905/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

CAS roles have been modified effective December 16, 2001. Please check your SDI profiles to see if they need to be revised. For information on CAS roles, enter HELP ROLES at an arrow prompt or use the CAS Roles thesaurus (/RL field) in this file.

=> s l3

L4 67 L3

=> s l4 and ((corticotropin releasing factor) or 'CRF')

11140 CORTICOTROPIN

70928 RELEASING

731932 FACTOR

4438 CORTICOTROPIN RELEASING FACTOR

(CORTICOTROPIN (W) RELEASING (W) FACTOR)

6604 'CRF'

L5 0 L4 AND ((CORTICOTROPIN RELEASING FACTOR) OR 'CRF')

=> s l4 and cortico?

70864 CORTICO?

L6 0 L4 AND CORTICO?

=> s l4 and (anxiety or depression or stress or immune or anorexia or addiction or inflammatory or cardiovascular)

8718 ANXIETY

62208 DEPRESSION

367768 STRESS

138445 IMMUNE

4522 ANOREXIA

3580 ADDICTION

92715 INFLAMMATORY

09/ 930,782

52360 CARDIOVASCULAR

L7 2 L4 AND (ANXIETY OR DEPRESSION OR STRESS OR IMMUNE OR ANOREXIA
OR ADDICTION OR INFLAMMATORY OR CARDIOVASCULAR)

=> d 17 1- ibib abs hitstr

YOU HAVE REQUESTED DATA FROM 2 ANSWERS - CONTINUE? Y/(N):y

L7 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1998:392921 CAPLUS

DOCUMENT NUMBER: 129:67733

TITLE: Synthesis of some novel pyrazole derivatives as potential antiinflammatory agents with minimum ulcerogenic activity

AUTHOR(S): El-Hawash, Soad A. M.; El-Mallah, A. I.

CORPORATE SOURCE: Dep. Pharm. Chem., Fac. Pharm., Univ. Alexandria, Alexandria, Egypt

SOURCE: Pharmazie (1998), 53(6), 368-373

CODEN: PHARAT; ISSN: 0031-7144

PUBLISHER: Govi-Verlag Pharmazeutischer Verlag

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 129:67733

AB Several N-substituted amino-4-(3,4-dimethoxyphenyl)hydroxy-1H-pyrazole-1-carboxamides and -thiocarboxamides and a pyrazolo[1,5-a]-1,3,5-triazine were synthesized starting from 3,4-(MeO)2C6H3CH(CN)CO2Et. The new compds. were tested for in-vivo anti-inflammatory activity against carrageenan-induced rat paw edema. All compds. exhibited activity in the range of 23-65%. The most potent compds. were further evaluated for their ulcerogenic liability and acute toxicity. They were found to be less toxic and nearly devoid of ulcerogenic activity as compared to phenylbutazone and indometacin.

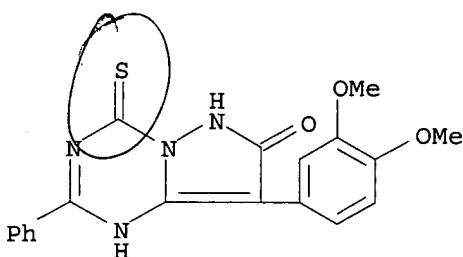
IT 209003-29-2P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)

(prepn. and anti-inflammatory activity of pyrazoles with min. ulcerogenic activity)

RN 209003-29-2 CAPLUS

CN Pyrazolo[1,5-a]-1,3,5-triazin-7(6H)-one, 8-(3,4-dimethoxyphenyl)-1,4-dihydro-2-phenyl-4-thioxo- (9CI) (CA INDEX NAME)



L7 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1998:1479 CAPLUS

DOCUMENT NUMBER: 128:61529

TITLE: Preparation and formulation of amide derivatives as pharmaceuticals

INVENTOR(S): Okamura, Takashi; Shoji, Yasuo; Shibutani, Tadao; Yasuda, Tsuneo; Iwamoto, Takeshi

PATENT ASSIGNEE(S): Otsuka Pharmaceutical Factory, Inc., Japan; Okamura, Takashi; Shoji, Yasuo; Shibutani, Tadao; Yasuda, Tsuneo; Iwamoto, Takeshi

SOURCE: PCT Int. Appl., 95 pp.

CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9746560	A1	19971211	WO 1997-JP1875	19970602
W: AU, CA, CN, JP, KR, US				
RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
CA 2257222	AA	19971211	CA 1997-2257222	19970602
AU 9729778	A1	19980105	AU 1997-29778	19970602
AU 716633	B2	20000302		
EP 915093	A1	19990512	EP 1997-924299	19970602
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				
CN 1221419	A	19990630	CN 1997-195345	19970602
CN 1066448	B	20010530		
US 6166016	A	20001226	US 1998-194727	19981202
KR 2000016395	A	20000325	KR 1998-709969	19981205
PRIORITY APPLN. INFO.:			JP 1996-144099	A 19960606
			JP 1997-73116	A 19970326
			WO 1997-JP1875	W 19970602
OTHER SOURCE(S):		MARPAT 128:61529		
GI				

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

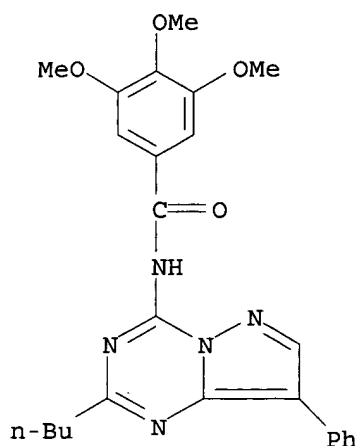
AB Amide derivs. I [ring A = benzene, naphthalene, pyridine or furan ring; when ring A is other than benzene ring, R1 = R2 = R3 = H; when ring A is a benzene ring, R1 - R3 = H, alkoxy, etc.; R4 = heterocycle such as lower alkyl-substituted thieno[3,2-d]pyrimidin-4-yl, optionally substituted pyrazolo[1,5-a]-1,3,5-triazin-4-yl, 6-substituted pyrazolo[3,4-d]pyrimidin-4-yl or 2-substituted purin-6-yl; and R5 = H, etc.] are prepd. I are analgesic, antiinflammatory, antimicrobial, hypoglycemic, hypolipemic, hypotensive, carcinostatic agents, etc. The title compd. II at 1 mg/kg showed analgesic activity in rats.

IT **200292-02-0P 200292-12-2P**
 RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (prepn. of amide derivs. as pharmaceuticals)

RN 200292-02-0 CAPLUS

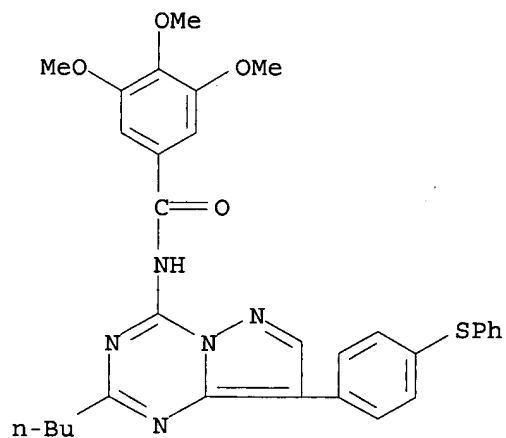
CN Benzamide, N-(2-butyl-8-phenylpyrazolo[1,5-a]-1,3,5-triazin-4-yl)-3,4,5-trimethoxy- (9CI) (CA INDEX NAME)

09/ 930,782



RN 200292-12-2 CAPLUS

CN Benzamide, N-[2-butyl-8-[4-(phenylthio)phenyl]pyrazolo[1,5-a]-1,3,5-triazin-4-yl]-3,4,5-trimethoxy- (9CI) (CA INDEX NAME)



=> d his

(FILE 'HOME' ENTERED AT 11:23:42 ON 06 SEP 2002)

FILE 'REGISTRY' ENTERED AT 11:23:50 ON 06 SEP 2002

L1 STRUCTURE UPLOADED

L2 14 S L1

L3 324 S L1 FUL

FILE 'CAPLUS' ENTERED AT 11:24:22 ON 06 SEP 2002

L4 67 S L3

L5 0 S L4 AND ((CORTICOTROPIN RELEASING FACTOR) OR 'CRF')

L6 0 S L4 AND CORTICO?

L7 2 S L4 AND (ANXIETY OR DEPRESSION OR STRESS OR IMMUNE OR ANOREXIA

=> s l4 and (gastrointestin? or alzheimer? or fertility or epilepsy or stroke?)

40330 GASTROINTESTIN?

23108 ALZHEIMER?

26357 FERTILITY

09/ 930,782

11374 EPILEPSY

17754 STROKE?

L8 2 L4 AND (GASTROINTESTIN? OR ALZHEIMER? OR FERTILITY OR EPILEPSY
OR STROKE?)

=> s l8 not l7

L9 2 L8 NOT L7

=> d l9 1- ibib abs hitstr

YOU HAVE REQUESTED DATA FROM 2 ANSWERS - CONTINUE? Y/(N):y

L9 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 2000:311637 CAPLUS

DOCUMENT NUMBER: 133:83844

TITLE: In-vivo sex differences in disposition and metabolism
of sulfoxide-containing drug BOF-4272 in rats

AUTHOR(S): Naito, Shinsaku; Nishimura, Masuhiro; Nogawa,
Hiroyuki; Yoshitsugu, Hiroki; Tamao, Yumi; Asano,
Satoru; Nomura, Yuuko

CORPORATE SOURCE: Naruto Research Institute, Otsuka Pharmaceutical
Factory, Inc., Tokushima, 772-8601, Japan

SOURCE: Pharmacy and Pharmacology Communications (2000), 6(4),
179-186

CODEN: PPCOFN; ISSN: 1460-8081

PUBLISHER: Royal Pharmaceutical Society of Great Britain

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Detailed sex differences in the disposition and metab. of BOF-4272, a
newly developed xanthine oxidase/xanthine dehydrogenase inhibitor, are
described on the basis of in-vivo studies in rats. In both male and
female rats, at all times after oral administration of 14C-labeled
BOF-4272, the highest level of radioactivity (except for radioactivity in
the **gastrointestinal** tract) was obsd. in the liver, then in the
kidney. Little radioactivity was detected in other tissues. After oral
administration the total 14C concn. profiles in the plasma and kidney were
almost identical in male and female rats whereas total 14C concns. in the
liver to 24 h were higher in female rats than in males. Levels of
BOF-4269 (the sulfide metabolite of BOF-4272) in the contents of the large
intestine to 24 h after the oral administration of BOF-4272 were higher in
female rats than in males. BOF-4269 was the main metabolite found in the
plasma in female rats after i.v. or oral administration. The area under
the plasma concn.-time curve (AUC0-t) for BOF-4272 after oral
administration was almost identical in female and male rats, whereas that
for BOF-4269 was 2.8 times higher in female rats than in males. These
findings suggest that differences between the disposition of 14C-labeled
BOF-4272 in rats of different sexes are mainly because of differences
between the metab. of BOF-4272 to BOF-4269 by the intestinal flora, the
elimination of BOF-4269 by the liver, or both.

IT 117411-21-9, BOF-4269 117411-36-6, BOF-4276
226925-31-1 226925-31-1D, glucuronides
226925-33-3 226925-35-5 226925-37-7
226925-37-7D, sulfates 260052-53-7D, sulfates
260052-54-8

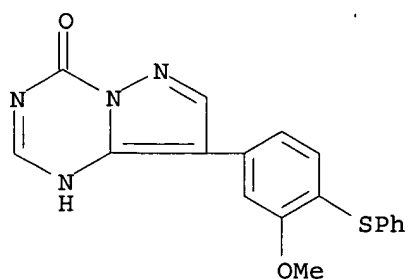
RL: BPR (Biological process); BSU (Biological study, unclassified); MFM
(Metabolic formation); BIOL (Biological study); FORM (Formation,
nonpreparative); PROC (Process)

(in-vivo sex differences in disposition and metab. of BOF-4272 in rats)

RN 117411-21-9 CAPLUS

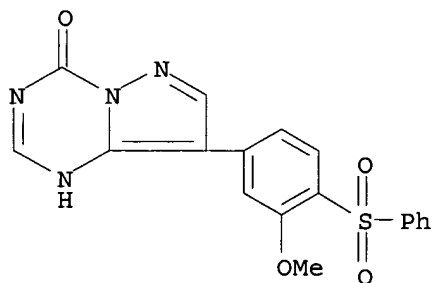
CN Pyrazolo[1,5-a]-1,3,5-triazin-4(1H)-one, 8-[3-methoxy-4-
(phenylthio)phenyl]- (9CI) (CA INDEX NAME)

09/ 930,782



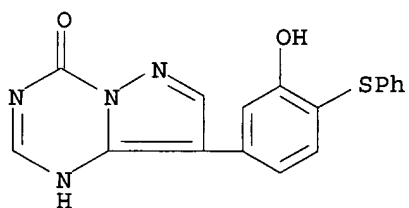
RN 117411-36-6 CAPLUS

CN Pyrazolo[1,5-a]-1,3,5-triazin-4(1H)-one, 8-[3-methoxy-4-(phenylsulfonyl)phenyl]- (9CI) (CA INDEX NAME)



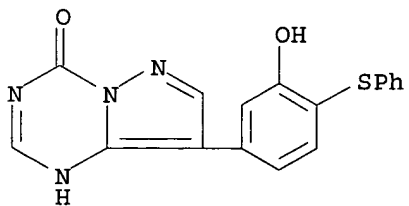
RN 226925-31-1 CAPLUS

CN Pyrazolo[1,5-a]-1,3,5-triazin-4(1H)-one, 8-[3-hydroxy-4-(phenylthio)phenyl]- (9CI) (CA INDEX NAME)



RN 226925-31-1 CAPLUS

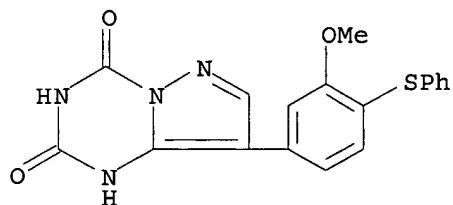
CN Pyrazolo[1,5-a]-1,3,5-triazin-4(1H)-one, 8-[3-hydroxy-4-(phenylthio)phenyl]- (9CI) (CA INDEX NAME)



RN 226925-33-3 CAPLUS

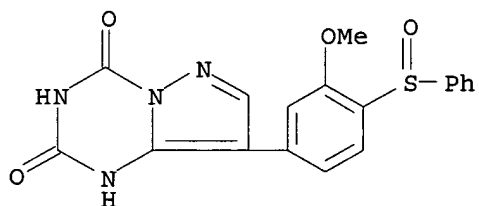
CN Pyrazolo[1,5-a]-1,3,5-triazine-2,4(1H,3H)-dione, 8-[3-methoxy-4-(phenylthio)phenyl]- (9CI) (CA INDEX NAME)

09/ 930,782



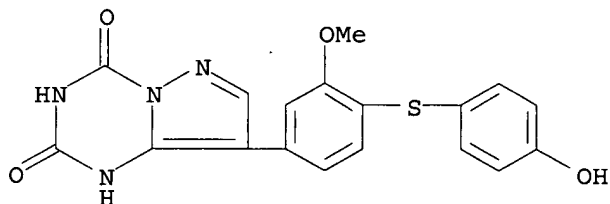
RN 226925-35-5 CAPLUS

CN Pyrazolo[1,5-a]-1,3,5-triazine-2,4(1H,3H)-dione, 8-[3-methoxy-4-(phenylsulfinyl)phenyl]- (9CI) (CA INDEX NAME)



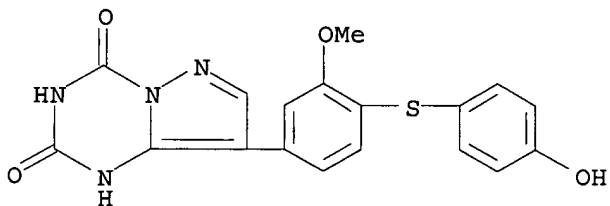
RN 226925-37-7 CAPLUS

CN Pyrazolo[1,5-a]-1,3,5-triazine-2,4(1H,3H)-dione, 8-[4-[(4-hydroxyphenyl)thio]-3-methoxyphenyl]- (9CI) (CA INDEX NAME)



RN 226925-37-7 CAPLUS

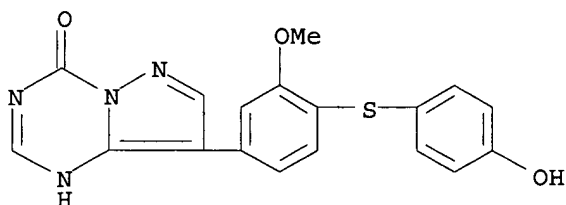
CN Pyrazolo[1,5-a]-1,3,5-triazine-2,4(1H,3H)-dione, 8-[4-[(4-hydroxyphenyl)thio]-3-methoxyphenyl]- (9CI) (CA INDEX NAME)



RN 260052-53-7 CAPLUS

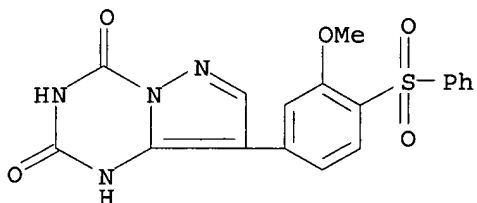
CN Pyrazolo[1,5-a]-1,3,5-triazine-4(1H)-one, 8-[4-[(4-hydroxyphenyl)thio]-3-methoxyphenyl]- (9CI) (CA INDEX NAME)

09/ 930,782



RN 260052-54-8 CAPLUS

CN Pyrazolo[1,5-a]-1,3,5-triazine-2,4(1H,3H)-dione, 8-[3-methoxy-4-(phenylsulfonyl)phenyl]- (9CI) (CA INDEX NAME)

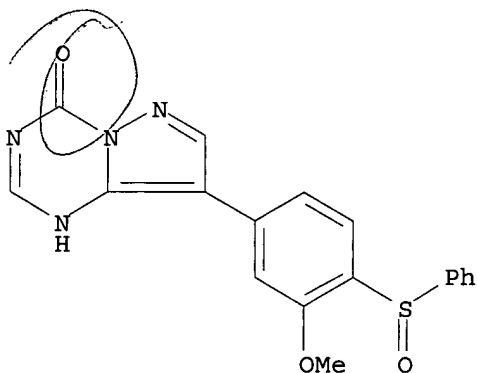


IT 142181-44-0, BOF-4272

RL: BPR (Biological process); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses) (in-vivo sex differences in disposition and metab. of BOF-4272' in rats)

RN 142181-44-0 CAPLUS

CN Pyrazolo[1,5-a]-1,3,5-triazin-4(1H)-one, 8-[3-methoxy-4-(phenylsulfinyl)phenyl]-, sodium salt (9CI) (CA INDEX NAME)



● Na

REFERENCE COUNT: 15 THERE ARE 15 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1995:637803 CAPLUS

DOCUMENT NUMBER: 123:40823

TITLE: Improvement of **gastrointestinal** absorption of a new oral xanthine oxidase/dehydrogenase inhibitor, sodium (.+-.)-8-(3-methoxy-4-phenylsulfinylphenyl)-pyrazolo[1,5-a]-1,3,5-triazine-4-olate monohydrate, in rats and dogs

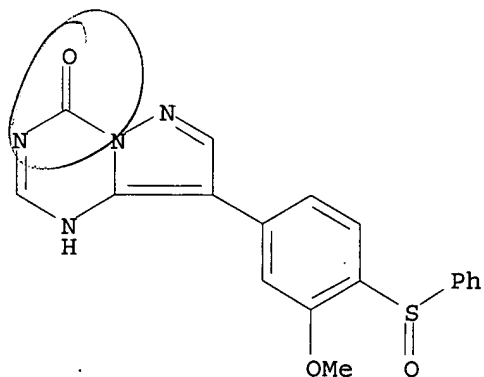
AUTHOR(S): Terao, Toshimitsu; Hatsubara, Takeshi; Kaga, Junji;

Kobatake, Hideki; Naito, Shinsaku
 CORPORATE SOURCE: Formulation Res. Inst., Otsuka Pharmaceutical Factory, Inc., Naruto, 772, Japan
 SOURCE: Yakuzai Gaku (1995), 55(1), 36-43
 CODEN: YAKUA2; ISSN: 0372-7629
 PUBLISHER: Nippon Yakuzai Gakkai
 DOCUMENT TYPE: Journal
 LANGUAGE: Japanese

AB Sodium (+-)-8-(3-methoxy-4-phenylsulfinylphenyl)pyrazolo[1,5-a]-1,3,5-triazine-4-olate monohydrate (BOF-4272) is a new synthetic inhibitor of xanthine oxidase/dehydrogenase. A preliminary rat expt. showed that the abs. bioavailability of BOF-4272 in oral administration was as low as 10%. In this study, taking the physicochem. properties of BOF-4272 into consideration, the absorption mechanism was investigated by the loop technique and the in situ loop technique using rats intestines. These results indicate that BOF-4272 is an acidic drug which is absorbed by passive transport. It was concluded that the low bioavailability of this compd. was due to not only the low permeability but also the low absorption rate attributable to the formation of ionized mols. in the **gastrointestinal** tract. The bioavailability was improved remarkably by the addn. of an enhancer (sugar ester) and by increasing the drug concn. in the small intestine.

IT 142181-44-0, BOF-4272
 RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
 (improvement of **gastrointestinal** absorption of a new oral xanthine oxidase/dehydrogenase inhibitor BOF-4272 in rats and dogs)

RN 142181-44-0 CAPLUS
 CN Pyrazolo[1,5-a]-1,3,5-triazin-4(1H)-one, 8-[3-methoxy-4-(phenylsulfinyl)phenyl]-, sodium salt (9CI) (CA INDEX NAME)



● Na

=> d his

(FILE 'HOME' ENTERED AT 11:23:42 ON 06 SEP 2002)

FILE 'REGISTRY' ENTERED AT 11:23:50 ON 06 SEP 2002

L1 STRUCTURE UPLOADED

L2 14 S L1

L3 324 S L1 FUL

FILE 'CAPLUS' ENTERED AT 11:24:22 ON 06 SEP 2002

09/ 930,782

L4 67 S L3
L5 0 S L4 AND ((CORTICOTROPIN RELEASING FACTOR) OR 'CRF')
L6 0 S L4 AND CORTICO?
L7 2 S L4 AND (ANXIETY OR DEPRESSION OR STRESS OR IMMUNE OR ANOREXIA
L8 2 S L4 AND (GASTROINTESTIN? OR ALZHEIMER? OR FERTILITY OR EPILEPS
L9 2 S L8 NOT L7

=> log y

COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
50.02	190.51

FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE	TOTAL
ENTRY	SESSION
-2.48	-2.48

CA SUBSCRIBER PRICE

STN INTERNATIONAL LOGOFF AT 11:29:22 ON 06 SEP 2002